# **Complete Summary**

#### **TITLE**

Hepatitis C: percentage of patients aged 18 years and older with a diagnosis of chronic hepatitis C who are receiving antiviral treatment for whom quantitative HCV RNA testing was performed within 6 months prior to initiation of antiviral treatment.

## SOURCE(S)

American Gastroenterological Association Institute, Physician Consortium for Performance Improvement®. Hepatitis C physician performance measurement set. Chicago (IL): American Medical Association (AMA); 2008 Jun. 42 p. [4 references]

## **Measure Domain**

#### PRIMARY MEASURE DOMAIN

**Process** 

The validity of measures depends on how they are built. By examining the key building blocks of a measure, you can assess its validity for your purpose. For more information, visit the Measure Validity page.

## SECONDARY MEASURE DOMAIN

Does not apply to this measure

## **Brief Abstract**

#### **DESCRIPTION**

This measure is used to assess the percentage of patients aged 18 years and older with a diagnosis of chronic hepatitis C who are receiving antiviral treatment for whom quantitative hepatitis C virus (HCV) ribonucleic acid (RNA) testing was performed within 6 months prior to initiation of antiviral treatment.

This measure is paired with <u>Hepatitis C: percentage of patients aged 18 years and older with a diagnosis of chronic hepatitis C who are receiving antiviral treatment for whom HCV genotype testing was performed prior to initiation of antiviral treatment. Implementers of this measure should not use this measure without the HCV genotype testing prior to treatment measure.</u>

#### **RATIONALE**

Establish baseline level against which to monitor virologic response and indicate likelihood of response. The clinical utility of serial hepatitis C virus (HCV) viral levels in a patient is predicated on continued use of the same specific quantitative assay that was used in the initial determination of the viral level. While there is little correlation between disease severity or disease progression with the absolute level of hepatitis C virus (HCV) ribonucleic acid (RNA), quantitative determination of the HCV level provides important information on the likelihood of response to treatment in patients undergoing antiviral therapy.\* (National Institutes of Health [NIH])

\*The following clinical recommendation statements are quoted <u>verbatim</u> from the referenced clinical guidelines and represent the evidence base for the measure:

HCV RNA testing should be performed in patients with a positive anti-HCV test, patients for whom antiviral treatment is being considered, using a quantitative assay, patients with unexplained liver disease whose anti-HCV test is negative and patients who are immunocompromised or suspected of having acute HCV infection. (American Association for the Study of Liver Diseases [AASLD])

All candidates for antiviral therapy should be tested for HCV RNA with a quantitative amplification assay, which provides both a baseline level against which to monitor virologic response and a prognostic indicator of the likelihood of response. (American Gastroenterological Association [AGA])

The diagnosis of chronic hepatitis C infection is often suggested by abnormalities in Alanine transaminase (ALT) levels and is established by enzyme immunoassay (EIA) followed by confirmatory determination of HCV RNA. (NIH)

#### PRIMARY CLINICAL COMPONENT

Chronic hepatitis C virus (HCV); quantitative HCV; ribonucleic acid (RNA) testing

#### **DENOMINATOR DESCRIPTION**

All patients aged 18 years and older with a diagnosis of chronic hepatitis C who are receiving antiviral treatment (see the related "Denominator Inclusions/Exclusions" field in the Complete Summary)

## NUMERATOR DESCRIPTION

Patients for whom quantitative hepatitis C virus (HCV) ribonucleic acid (RNA) testing was performed within 6 months prior to initiation of antiviral treatment

## **Evidence Supporting the Measure**

#### **EVIDENCE SUPPORTING THE CRITERION OF QUALITY**

- A clinical practice guideline or other peer-reviewed synthesis of the clinical evidence
- One or more research studies published in a National Library of Medicine (NLM) indexed, peer-reviewed journal

## **NATIONAL GUIDELINE CLEARINGHOUSE LINK**

• Diagnosis, management, and treatment of hepatitis C.

# **Evidence Supporting Need for the Measure**

#### **NEED FOR THE MEASURE**

Unspecified

# **State of Use of the Measure**

#### STATE OF USE

Current routine use

# **CURRENT USE**

Internal quality improvement National reporting

# **Application of Measure in its Current Use**

#### **CARE SETTING**

Ambulatory Care Physician Group Practices/Clinics

## PROFESSIONALS RESPONSIBLE FOR HEALTH CARE

Physicians

## LOWEST LEVEL OF HEALTH CARE DELIVERY ADDRESSED

**Individual Clinicians** 

# **TARGET POPULATION AGE**

Age greater than or equal to 18 years

# **TARGET POPULATION GENDER**

Either male or female

## STRATIFICATION BY VULNERABLE POPULATIONS

Unspecified

# **Characteristics of the Primary Clinical Component**

# INCIDENCE/PREVALENCE

Unspecified

## **ASSOCIATION WITH VULNERABLE POPULATIONS**

Unspecified

## **BURDEN OF ILLNESS**

Unspecified

## **UTILIZATION**

Unspecified

## COSTS

Unspecified

**Institute of Medicine National Healthcare Quality Report Categories** 

# **IOM CARE NEED**

Living with Illness

## **IOM DOMAIN**

Effectiveness

# **Data Collection for the Measure**

## **CASE FINDING**

Users of care only

# **DESCRIPTION OF CASE FINDING**

All patients aged 18 years and older with a diagnosis of chronic hepatitis C who are receiving antiviral treatment

## **DENOMINATOR SAMPLING FRAME**

Patients associated with provider

## **DENOMINATOR INCLUSIONS/EXCLUSIONS**

## **Inclusions**

All patients aged 18 years and older with a diagnosis of chronic hepatitis C who are receiving antiviral treatment

#### **Exclusions**

Documentation of medical reason(s) for not performing hepatitis C virus (HCV) ribonucleic acid (RNA) within 6 months prior to initiation of antiviral treatment

#### RELATIONSHIP OF DENOMINATOR TO NUMERATOR

All cases in the denominator are equally eligible to appear in the numerator

## **DENOMINATOR (INDEX) EVENT**

Clinical Condition Encounter Therapeutic Intervention

## **DENOMINATOR TIME WINDOW**

Time window is a single point in time

## **NUMERATOR INCLUSIONS/EXCLUSIONS**

#### **Inclusions**

Patients for whom quantitative hepatitis C virus (HCV) ribonucleic acid (RNA) testing was performed within 6 months prior to initiation of antiviral treatment

#### **Exclusions**

None

# MEASURE RESULTS UNDER CONTROL OF HEALTH CARE PROFESSIONALS, ORGANIZATIONS AND/OR POLICYMAKERS

The measure results are somewhat or substantially under the control of the health care professionals, organizations and/or policymakers to whom the measure applies.

## **NUMERATOR TIME WINDOW**

Fixed time period

## **DATA SOURCE**

Administrative data Medical record

## LEVEL OF DETERMINATION OF QUALITY

Individual Case

## **PRE-EXISTING INSTRUMENT USED**

Unspecified

# **Computation of the Measure**

#### **SCORING**

Rate

## **INTERPRETATION OF SCORE**

Better quality is associated with a higher score

## **ALLOWANCE FOR PATIENT FACTORS**

Unspecified

## STANDARD OF COMPARISON

Internal time comparison

# **Evaluation of Measure Properties**

## **EXTENT OF MEASURE TESTING**

Unspecified

# **Identifying Information**

## **ORIGINAL TITLE**

Measure #2: hepatitis C ribonucleic acid (RNA) testing before initiating treatment.

## **MEASURE COLLECTION**

The Physician Consortium for Performance Improvement® Measurement Sets

## **MEASURE SET NAME**

Hepatitis C Physician Performance Measurement Set

#### **SUBMITTER**

American Medical Association on behalf of the American Gastroenterological Association Institute and Physician Consortium for Performance Improvement®

## **DEVELOPER**

American Gastroenterological Association Institute Physician Consortium for Performance Improvement®

# **FUNDING SOURCE(S)**

Unspecified

#### COMPOSITION OF THE GROUP THAT DEVELOPED THE MEASURE

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#### FINANCIAL DISCLOSURES/OTHER POTENTIAL CONFLICTS OF INTEREST

Conflicts, if any, are disclosed in accordance with the Physician Consortium for Performance Improvement® conflict of interest policy.

#### **ENDORSER**

National Quality Forum

#### **INCLUDED IN**

Ambulatory Care Quality Alliance Physician Quality Reporting Initiative

## **ADAPTATION**

Measure was not adapted from another source.

## **RELEASE DATE**

2006 Dec

## **REVISION DATE**

2008 Jun

#### **MEASURE STATUS**

This is the current release of the measure.

## SOURCE(S)

American Gastroenterological Association Institute, Physician Consortium for Performance Improvement®. Hepatitis C physician performance measurement set. Chicago (IL): American Medical Association (AMA); 2008 Jun. 42 p. [4 references]

#### MEASURE AVAILABILITY

The individual measure, "Measure #2: Hepatitis C Ribonucleic Acid (RNA) Testing Before Initiating Treatment," is published in "Hepatitis C Physician Performance Measurement Set." This document and technical specifications are available in Portable Document Format (PDF) from the American Medical Association (AMA)-convened Physician Consortium for Performance Improvement® Web site: www.physicianconsortium.org.

For further information, please contact AMA staff by e-mail at <a href="mailto:cgi@ama-assn.org">cgi@ama-assn.org</a>.

## **NQMC STATUS**

This NQMC summary was completed by ECRI Institute on February 27, 2009. The information was verified by the measure developer on May 21, 2009.

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